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(54) **METHODS AND SYSTEMS FOR EVALUATING BONE LESIONS**

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A61B 6/00	(2006.01)
A61B 5/00	(2006.01)

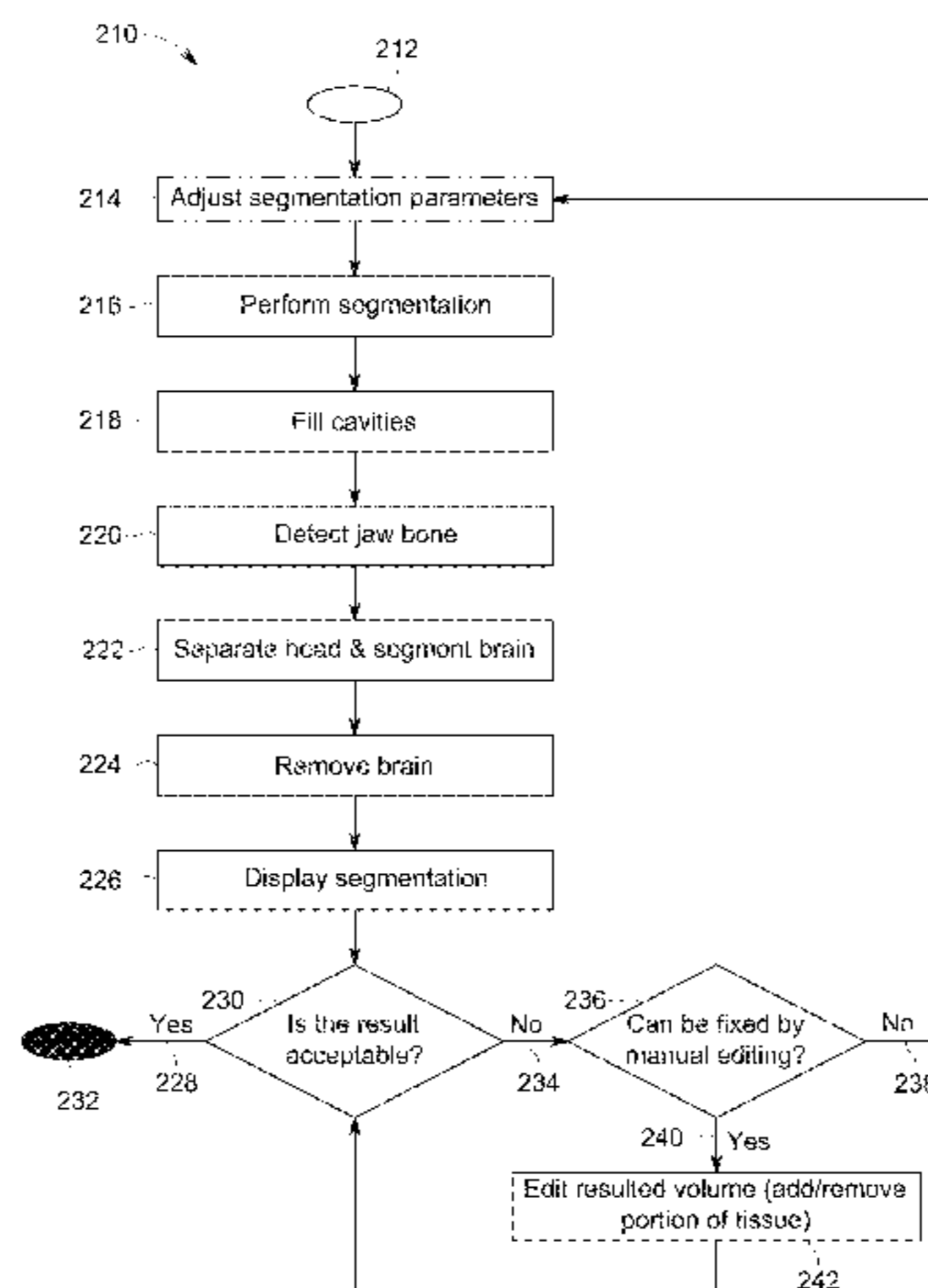
(57) **ABSTRACT**

Methods and systems for evaluating bone lesions include accessing a first dataset acquired from a patient with a first imaging modality and a second dataset acquired from the patient with a second imaging modality. A segmentation is performed on the first dataset to identify a subset of the first dataset corresponding to a skeletal structure of the patient and a patient skeletal metric representing a total bone volume of the patient is automatically calculated from the subset of the first dataset. The methods and systems further include detection of at least one lesion in the second dataset, classification of the at least one lesion as a bone or non-bone lesion, automatic calculation of a bone lesion metric based on the classification, and calculation of a lesion burden as a ratio of the bone lesion metric and the patient skeletal metric.

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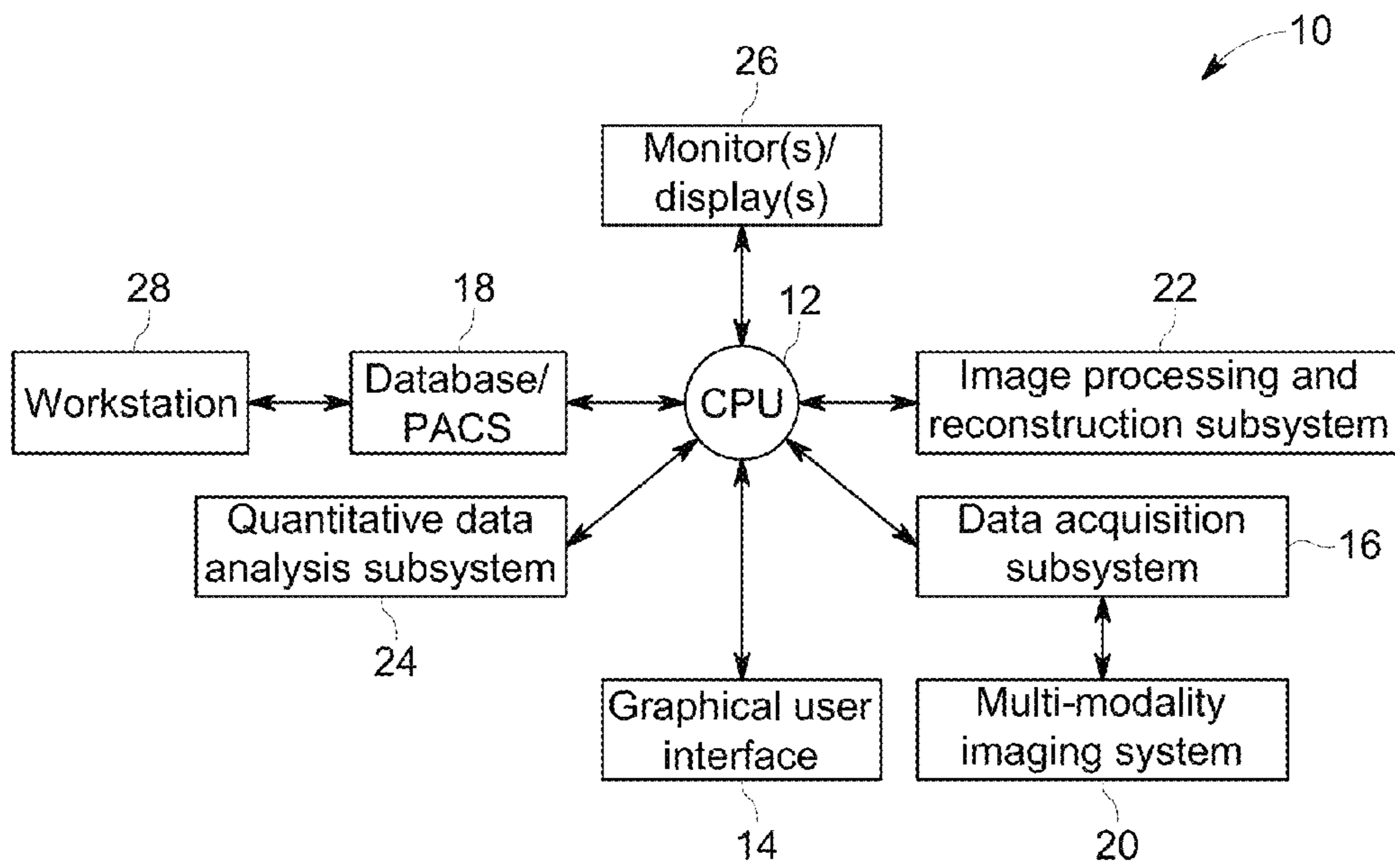
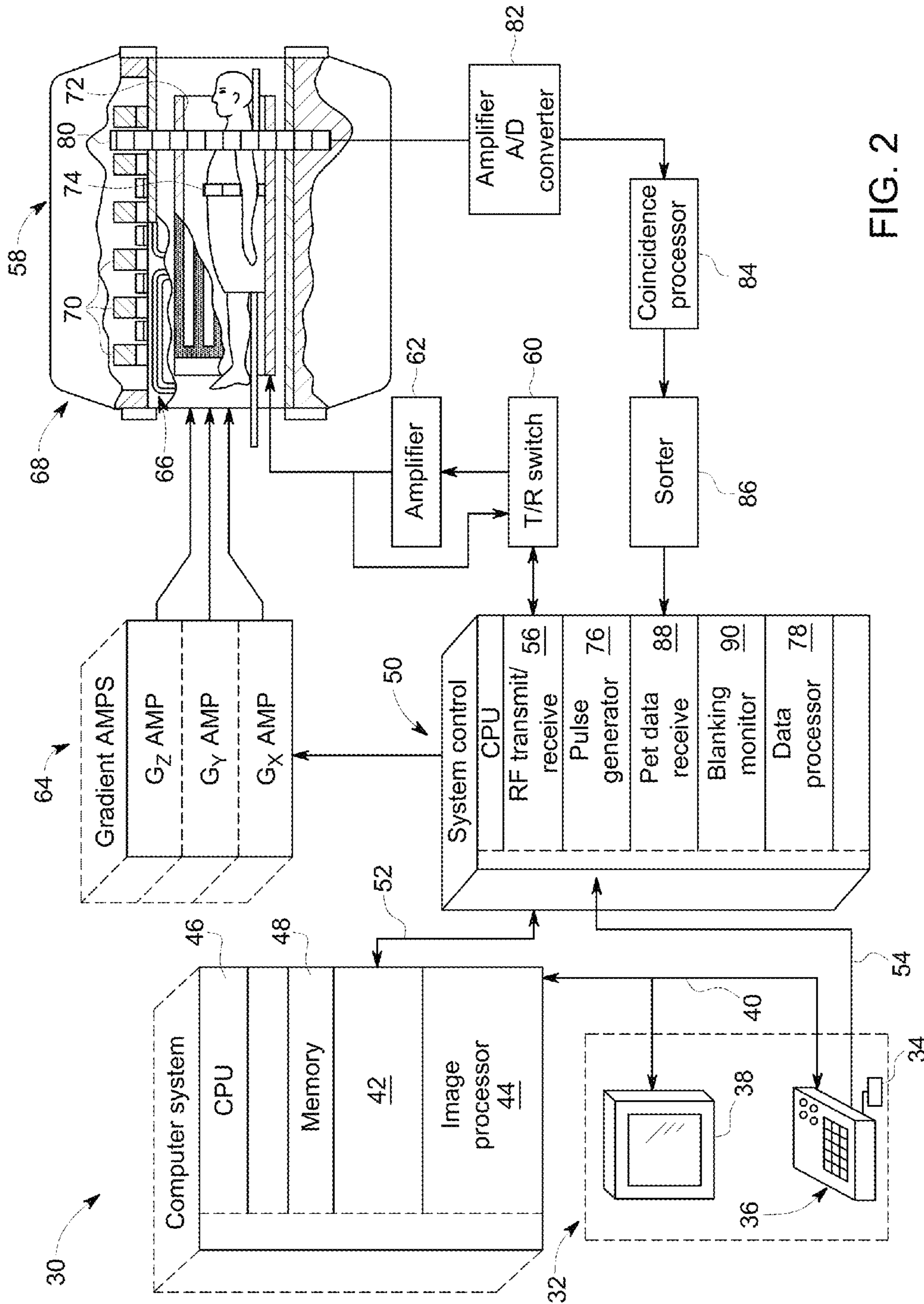


FIG. 1



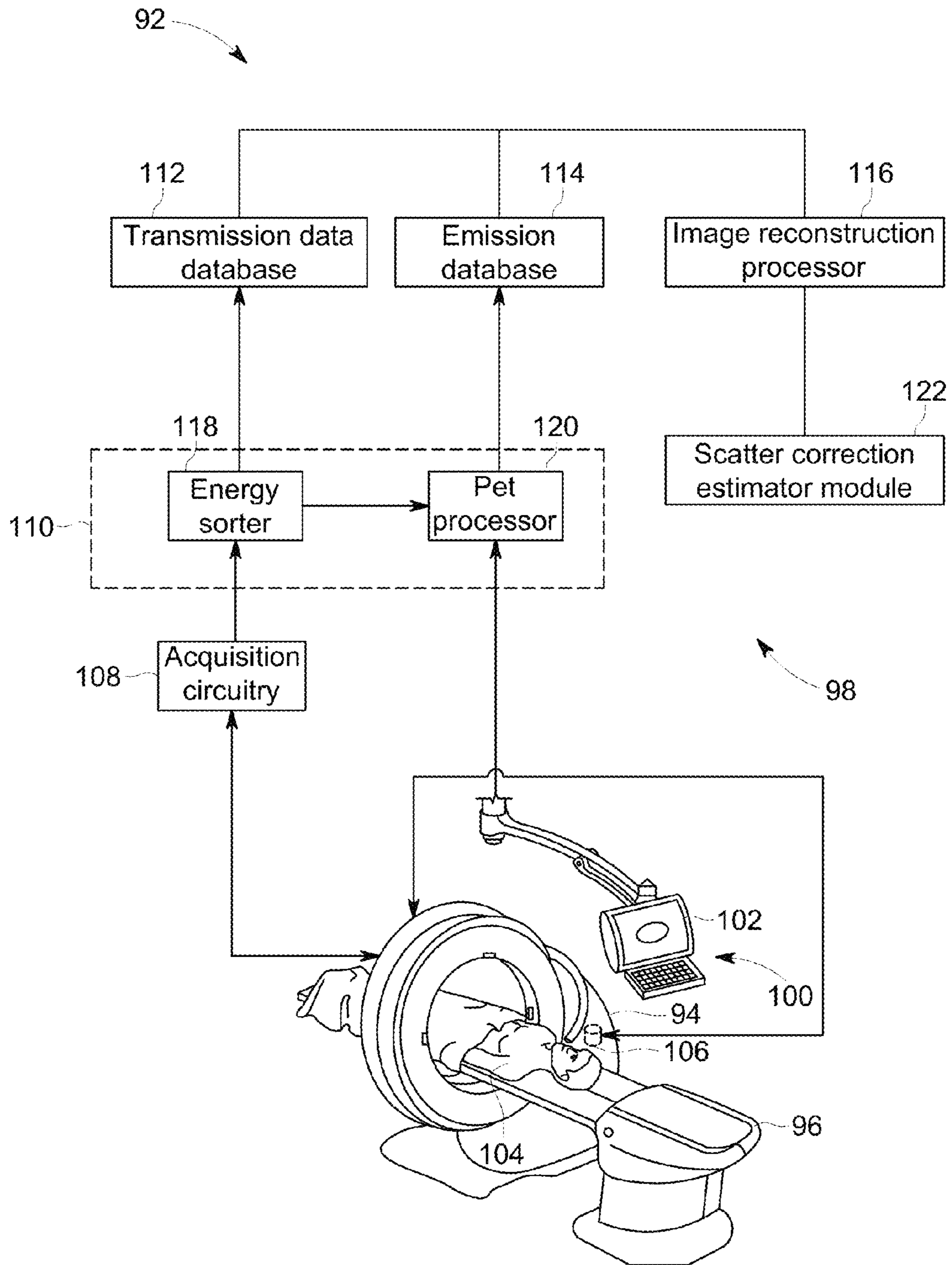


FIG. 3

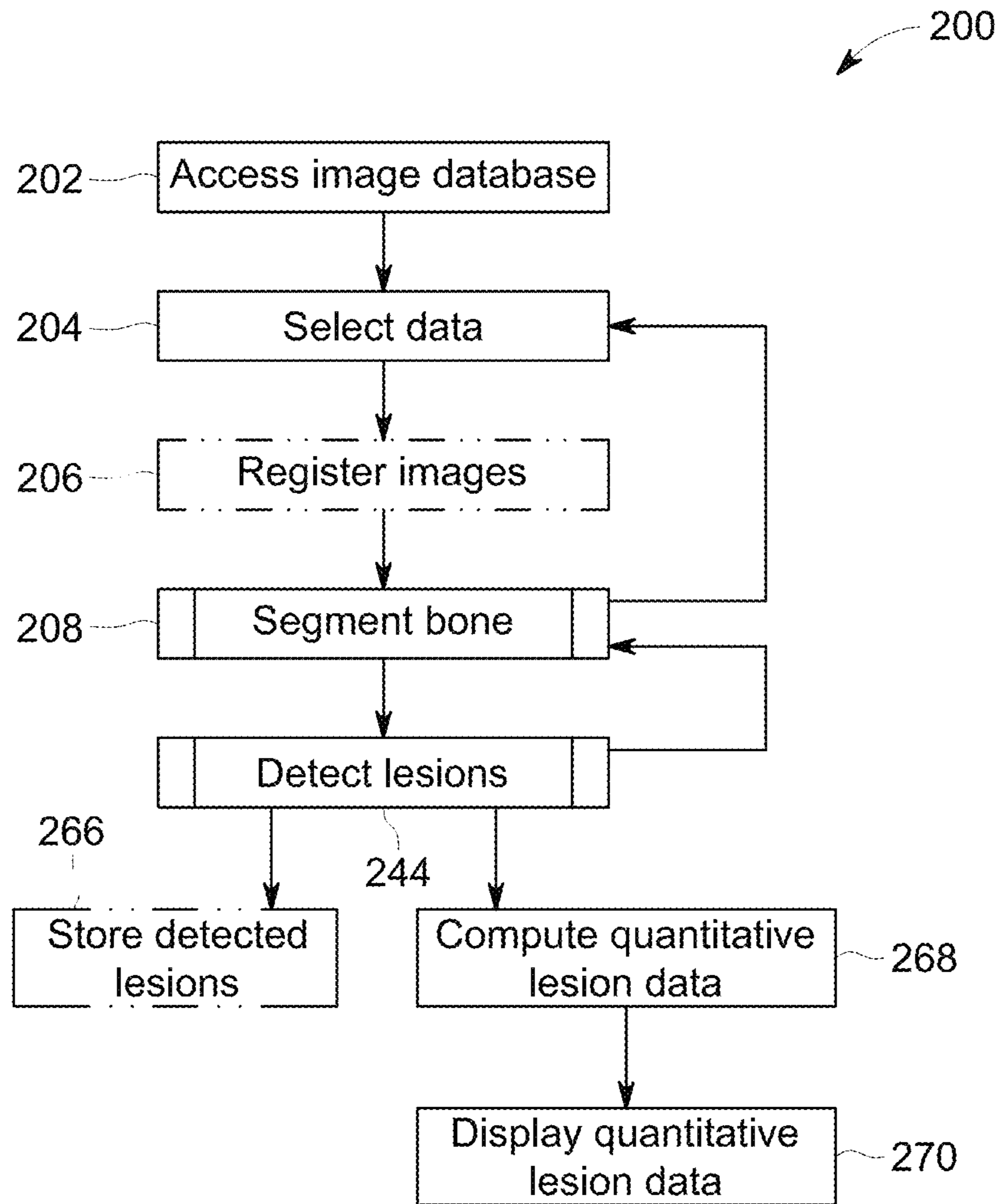


FIG. 4

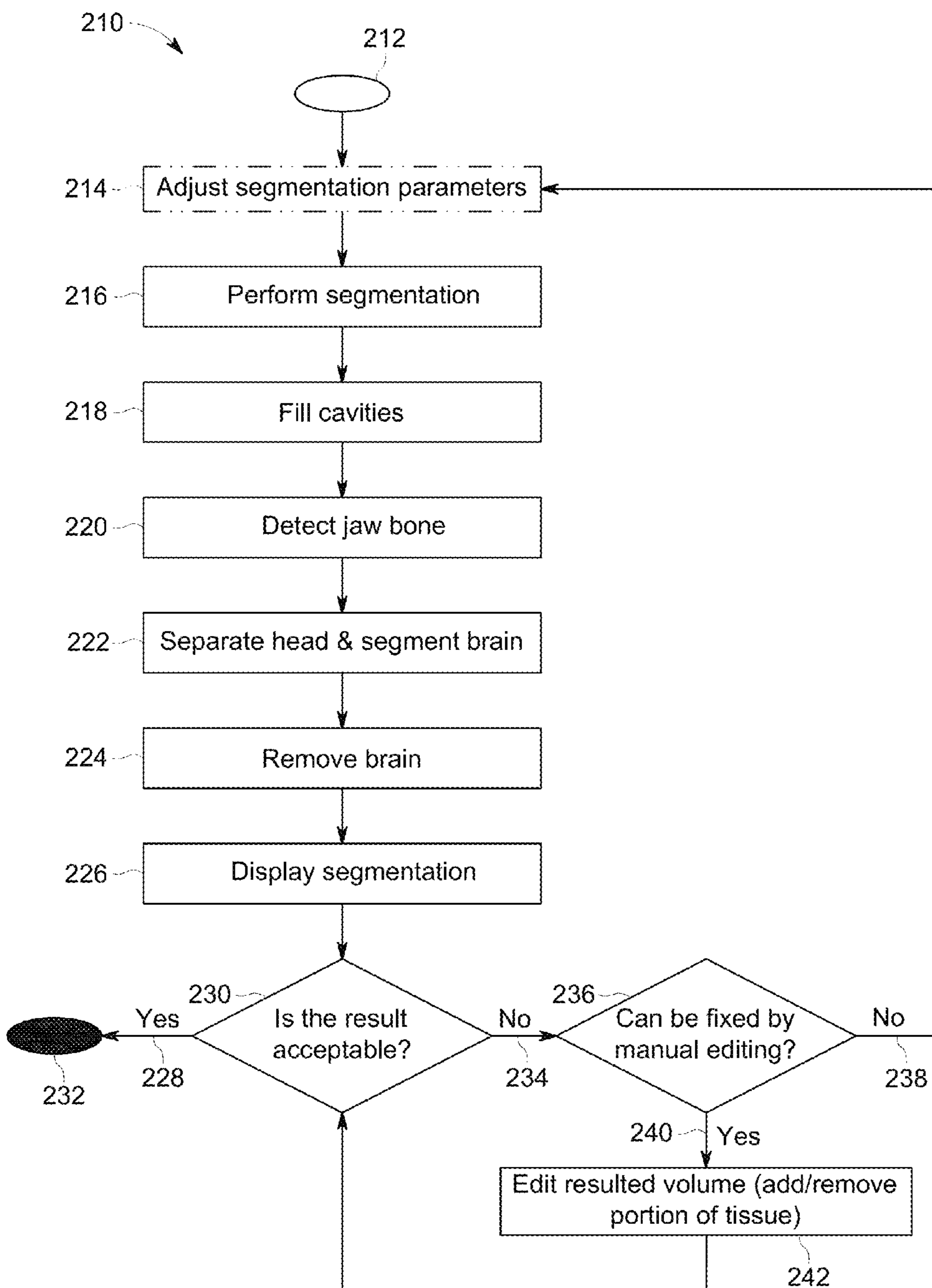


FIG. 5

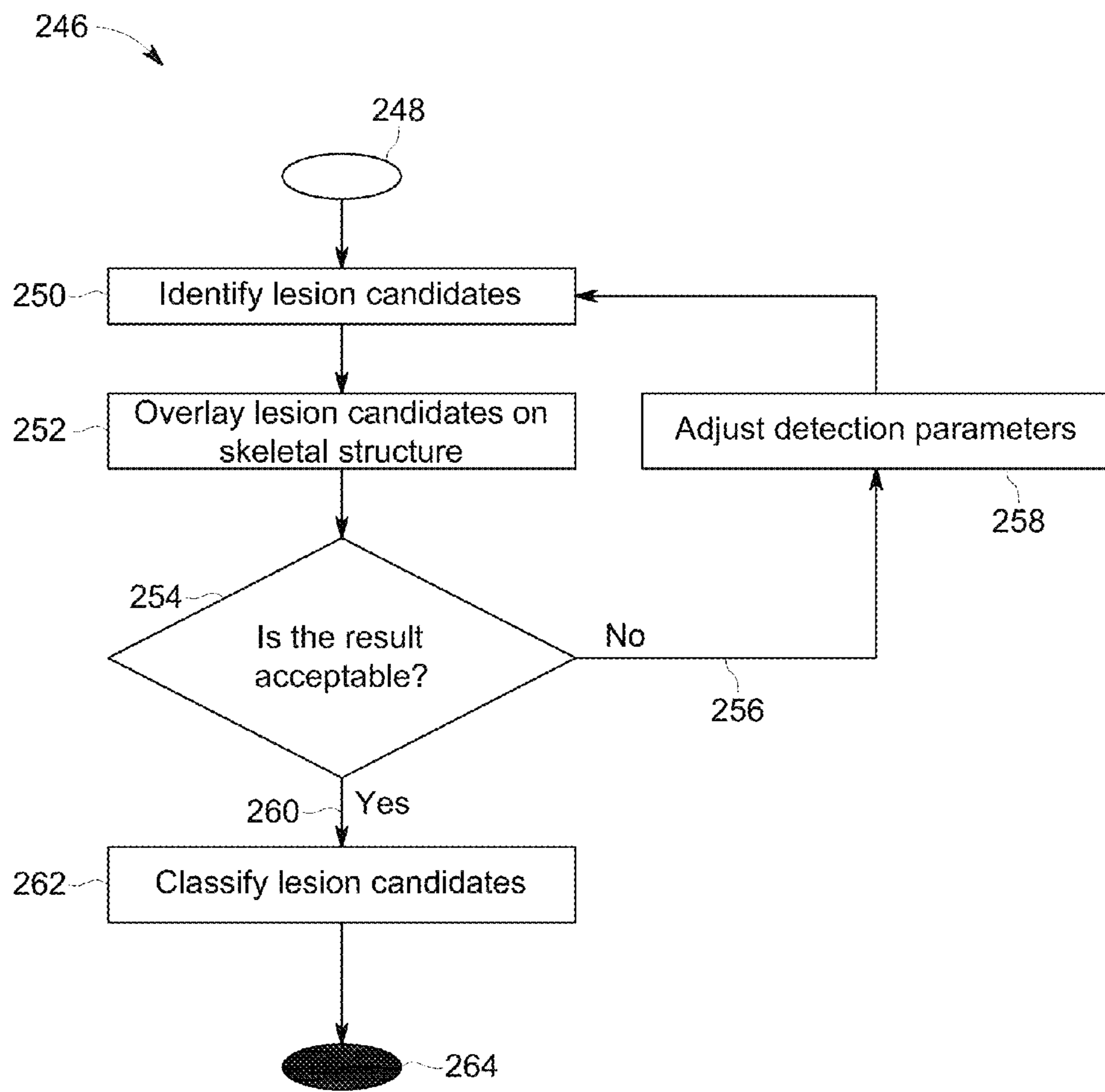
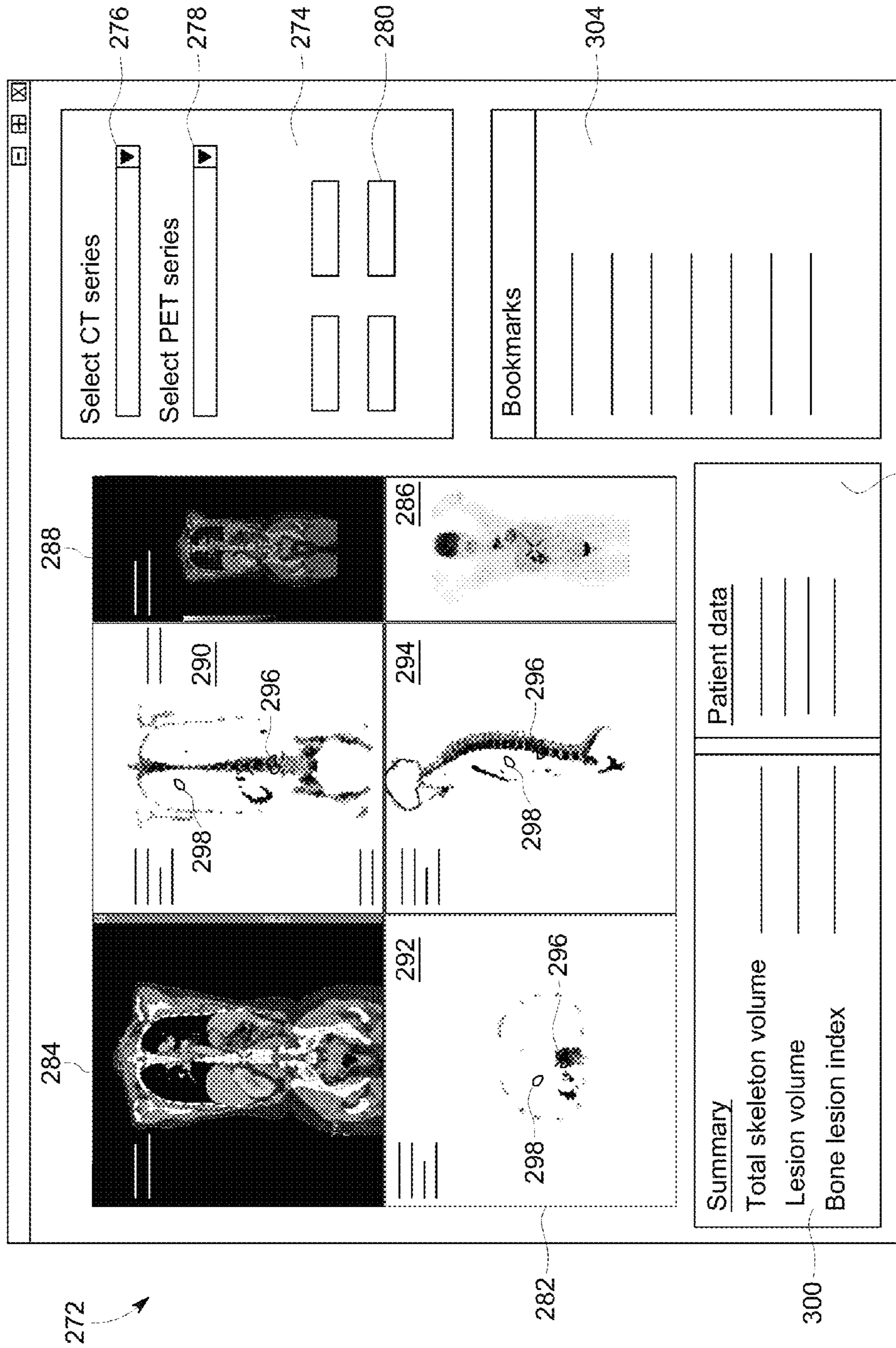


FIG. 6



302

FIG. 7

1

**METHODS AND SYSTEMS FOR
EVALUATING BONE LESIONS**

BACKGROUND OF THE INVENTION

Embodiments of the invention relate generally to medical diagnostics and treatment assessment and, more particularly, to methods and systems for evaluating bone lesions using multi-modality image data.

It is not uncommon for a single patient to undergo a multitude of medical imaging exams, whether in a single doctor's visit, in a hospital stay, or even over the course of a lifetime. This is particularly likely when a patient undergoes a series of "tests" and scans to investigate a recently onset or previously undetected condition. It is increasingly common for a patient to be subjected to multiple scans across multiple medical imaging modalities because each exam can provide different pieces of information. For example, during a single doctor's visit or hospital stay, a magnetic resonance (MR) imaging system, an x-ray imaging system, or a computed tomography (CT) imaging system can be used to acquire images that provide anatomical information, while a positron emission tomography (PET) imaging system or functional MRI can be used to acquire images that provide functional information. The anatomical information providing insight into the anatomical makeup of the patient and the functional information providing insight into the functionality of a given anatomical structure, especially when subjected to a stimulus. Moreover, the combination of anatomical and functional information is not only advantageous in detecting a new pathology or abnormality, but the respective images, when taken over the course of an illness, for example, may show growth of lesions, responses to treatments, and disease progression. To assist in the analysis of anatomical and functional information, programs have been constructed that register an anatomical and a functional image thereby showing, in a single image, both anatomical and functional information.

Many clinical applications analyze 2D or 3D image data to perform and capture quantitative analytics. These include detection and sizing of lung nodules (Advanced Lung Analysis), quantification of vessel curvature, diameter, and tone (Advanced Vessel Analysis), cardiac vascular and function applications, navigating of the colon for detection of polyps (CT colonography), detection and sizing of lesions, and the like. Dedicated CT, MR, PET and nuclear medicine applications have been designed to output quantitative analytics from regions of interest (intensity, density (HU), standard uptake value (SUV), distances, volumes, growth rates, pattern and/or texture recognition, functional information, etc.) to help in the diagnosis and management of patients.

Quantification of bone lesions using medical images is an important aspect of clinical diagnostics and therapy. Information related to the quantity or overall volume of bone lesions detected in a patient may be used by medical practitioners to select the best course of treatment for a patient and to monitor treatment efficiency and collect relevant research data.

Conventional methods of quantifying bone lesions in a patient utilize a reference value as an estimate of the volume of a particular patient's skeletal structure. Because this reference value is a quantitative value selected from tabular reference data based on the demographics (e.g., age, sex, height, etc.) of the patient, the reference value may not accurately reflect the actual skeletal volume of the patient.

Lesions within the skeletal structure of the patient are detected by first manually segmenting the skeletal structure from image data acquired from the patient and manually identifying lesions within the segmentation. The manual seg-

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mentation of the skeletal structural and manual detection of lesions is very challenging due to the complexity of the skeletal structure. For example, the anatomy and composition of bone material can vary significantly among patients, which can lead to significant inter-operator variability.

Further, the segmentation of the skeletal structure typically includes the brain, which can cause inaccurate quantitative analytics of the skeletal structure, and in turn affects a practitioner's ability to efficiently and accurately acquire measurements and data to assess the patient's condition.

Therefore, it would be desirable to have a system and method of quantifying bone lesions that overcomes the aforementioned drawbacks.

BRIEF DESCRIPTION OF THE INVENTION

In accordance with one aspect of the present invention, a non-transitory computer readable storage medium has stored thereon a computer program and represents a set of instructions that when executed by the computer causes the computer to access a first set of data of a first data type and access a second set of data of a second data type. The first set of data is acquired from a patient with a first imaging modality, and the second set of data is acquired from the patient with a second imaging modality. The set of instructions also causes the computer to perform a segmentation on the first set of data to identify a subset of the first set of data corresponding to a skeletal structure of the patient. Further, the set of instructions causes the computer to automatically calculate a patient skeletal metric from the subset of the first set of data, the patient skeletal metric representing a total bone volume of the patient. In addition, the set of instructions causes the computer to detect at least one lesion in the second set of data, classify the at least one lesion as a one of a bone lesion and a non-bone lesion, and automatically calculate a bone lesion metric based on the classification. The set of instructions further causes the computer to calculate a lesion burden as a ratio of the bone lesion metric and the patient skeletal metric.

According to another aspect of the invention, a method includes accessing an anatomical image data set acquired from a patient and accessing a function image data set acquired from the patient. The method also includes identifying a subset of the anatomical image dataset corresponding to bone and calculating a skeletal volume from the subset of the anatomical image dataset. In addition, the method includes identifying a set of lesions from the function image dataset, comparing the set of lesions to the subset of the anatomical image dataset to identify at least one bone lesion within the set of lesions, automatically calculating a bone lesion burden based on the at least one bone lesion. Further, the method includes calculating a bone lesion index from the bone lesion burden and the skeletal volume.

In accordance with another aspect of the invention, a medical diagnostic tool includes an image acquisition system configured to acquire multi-modality image data, a first database having stored thereon a first image dataset acquired from a patient using a first imaging modality, a second database having stored thereon a second image dataset acquired from the patient using a second imaging modality, and a computer processor. The computer processor is programmed to access the first and second image datasets. The computer processor is further programmed to segment the first image dataset to define a set of skeletal data, identify a location of a lesion candidate in the second image set, compare the location of the lesion candidate to the set of skeletal data, and classify the lesion candidate as one of a bone lesion and a non-bone lesion based on the comparison to define a set of bone lesion data. In

addition, the computer processor is programmed to calculate a lesion burden based on a ratio of the set of bone lesion data and the set of skeletal data. The medical diagnostic tool also includes a graphical user interface (GUI) constructed to display an image generated from overlaying a first image corresponding to the set of skeletal data on a second image corresponding to the set of bone lesion data.

Various other features and advantages will be made apparent from the following detailed description and the drawings.

BRIEF DESCRIPTION OF THE DRAWINGS

The drawings illustrate preferred embodiments presently contemplated for carrying out the invention.

In the drawings:

FIG. 1 is a schematic diagram of a medical imaging system according to one embodiment of the invention.

FIG. 2 is a schematic block diagram of an exemplary multi-modality medical imaging system, useable with the medical imaging system of FIG. 1, which includes a PET imaging system and a MR imaging system.

FIG. 3 is a schematic block diagram of an exemplary multi-modality medical imaging system, useable with the medical imaging system of FIG. 1, which includes a PET imaging system and a CT imaging system.

FIG. 4 is a flowchart setting forth the steps of a technique for quantifying bone lesions using multi-modality image data according to one embodiment of the invention.

FIG. 5 is a flowchart setting forth the steps of a bone segmentation subroutine of the technique of FIG. 4 according to one embodiment of the invention.

FIG. 6 is a flowchart setting forth the steps of a lesion detection subroutine of the technique of FIG. 4 according to one embodiment of the invention.

FIG. 7 illustrates an exemplary visual representation of a graphical user interface (GUI) for displaying a visualization of data in accordance with one embodiment of the present invention.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

The operating environment of embodiments of the invention is described below with respect to a multi-modality imaging system that includes a positron emission tomography (PET) imaging system and one of an magnetic resonance (MR) imaging system and a sixty-four-slice, "third generation" computed tomography (CT) imaging system. However, it will be appreciated by those skilled in the art that the invention is equally applicable for use with other multi-slice configurations and with other CT imaging systems. In addition, while embodiments of the invention are described with respect to techniques for use with MR or CT imaging systems and PET imaging systems, one skilled in the art will recognize that the concepts set forth herein are not limited to CT and PET and can be applied to techniques used with other imaging modalities in both the medical field and non-medical field, such as, for example, an x-ray system, a single-photon emission computed tomography (SPECT) imaging system, or any combination thereof.

Referring now to FIG. 1, an exemplary imaging system 10 for use with embodiments of the present invention is shown. The imaging system 10 is not modality specific. In this regard, the system 10 includes a central processor (CPU) 12 that controls operation of the system 10. The imaging system 10 further includes a graphical user interface (GUI) 14 that allows an operator to prescribe a scan and interact with the

collected data and images. Data acquisition is controlled, via the CPU 12, by data acquisition subsystem 16. According to various embodiments, data acquisition subsystem 16 may access previously acquired data stored in database 18 or may acquire data directly from a multi-modality imaging subsystem 20. Multi-modality imaging subsystem 20 may be a combined PET/MR system (as shown in FIG. 2), a combined PET/CT system (as shown in FIG. 3), a combined SPECT/MR system, or a combined SPECT/CT system, as non-limiting examples.

Imaging system 10 further includes an image processing and reconstruction subsystem 22 that generates reconstructed images as well a quantitative data analysis subsystem 24 that derives quantitative data from data collected from data acquisition system 16. The reconstructed images generated by image processing and reconstruction subsystem 22 and the quantitative data computed or otherwise derived from quantitative data analysis subsystem 24 may be stored in database 18. Additionally, it is contemplated that database 18 may be more than one database and remotely located from the site of data acquisition.

System 10 further has one or more displays 26 to visually display images and quantitative data as set forth herein. One having skill in the art will appreciate that that system 10 may include other software, firmware, and hardware not specifically described to prescribe and execute a given scan, as well as processing the data for image reconstruction and quantitative analysis.

Additionally, it is contemplated that a dedicated workstation 28 having a computer, monitor(s), and operably connected to the one or more databases may be used such that a physician may analyze the image and quantitative data remote from the scanner. As such, it is understood that systems and methods described herein may be used remotely from the treatment facility at which the patient is scanned.

FIG. 2 illustrates an exemplary PET/MR imaging system 30 useable with multi-modality imaging subsystem 20. The operation of PET/MR imaging system 30 may be controlled from an operator console 32, which includes a keyboard or other input device 34, a control panel 36, and a display 38. The console 32 communicates through a link 40 with a separate computer system 42 that enable an operator to control the production and display of images on the display 38. The computer system 42 includes a number of modules, such as an image processor module 44, a CPU module 46, and a memory module 48. The computer system 42 may also be connected to permanent or back-up memory storage, a network, or may communicate with a separate system control 50 through link 52. The input device 34 can include a mouse, keyboard, track ball, touch activated screen, light wand, or any similar or equivalent input device, and may be used for interactive geometry prescription.

The system control 50 includes a set of modules in communication with one another and connected to the operator console 32 through link 54. It is through link 52 that the system control 50 receives commands to indicate the scan sequence or sequences that are to be performed. For MR data acquisition, an RF transmit/receive module 56 commands a scanner 58 to carry out the desired scan sequence, by sending instructions, commands, and/or requests describing the timing, strength, and shape of the RF pulses and pulse sequences to be produced, to correspond to the timing and length of the data acquisition window. In this regard, a transmit/receive switch 60 controls the flow of data via an amplifier 62 to scanner 58 from RF transmit module 56 and from scanner 58 to RF receive module 56. The system control 50 also connects

to a set of gradient amplifiers **64**, to indicate the timing and shape of the gradient pulses that are produced during the scan.

The gradient waveform instructions produced by system control **50** are sent to the gradient amplifier system **64** having Gx, Gy, and Gz amplifiers. Amplifiers **64** may be external of scanner **58** or system control **50**, or may be integrated therein. Each gradient amplifier excites a corresponding physical gradient coil in a gradient coil assembly **66** to produce the magnetic field gradients used for spatially encoding acquired signals. The gradient coil assembly **66** forms part of a magnet assembly **68**, which includes a polarizing magnet **70** and an RF coil assembly **72**, **74**. Alternatively, the gradient coils of gradient coil assembly **66** may be independent of the magnet assembly **68**. RF coil assembly may include a whole-body RF transmit coil **72** as shown, surface or parallel imaging coils **74**, or a combination of both. The coils **72**, **74** of the RF coil assembly may be configured for both transmitting and receiving, or for transmit-only or receive-only. A pulse generator **76** may generate PET data blanking signals synchronously with the production of the pulse sequences. These blanking signals may be generated on separate logic lines for subsequent data processing. The MR signals resulting from the excitation pulses, emitted by the excited nuclei in the patient, may be sensed by the whole body coil **72** or by separate receive coils, such as parallel coils or surface coils **74**, and are then transmitted to the RF transmit/receive module **56** via the transmit/receive switch **60**. The MR signals are demodulated, filtered, and digitized in a data processing section **78** of the system control **50**.

An MR scan is complete when one or more sets of raw k-space data has been acquired in the data processor **78**. This raw k-space data is reconstructed in data processor **78**, which operates to transform the data (through Fourier or other techniques) in image data. This image data is conveyed through link **52** to the computer system **42**, where it is stored in memory **48**. Alternatively, in some systems computer system **42** may assume the image data reconstruction and other functions of data processor **78**. In response to commands received from the operator console **32**, the image data stored in memory **48** may be archived in long term storage or may be further processed by the image processor **44** or CPU **46** and conveyed to the operator console **32** and presented on the display **38**.

In combined PET/MR imaging systems, PET data may be acquired simultaneously with the MR data acquisition described above. Thus, scanner **58** also contains a positron emission detector **80**, configured to detect gamma rays from positron annihilations emitted from a subject. Detector **80** preferably includes a plurality of scintillators and photovoltaics arranged about a gantry. Detector **80** may, however, be of any suitable construction for acquiring PET data. Gamma ray incidences detected by detector **80** are transformed, by the photovoltaics of the detector **80**, into electrical signals and are conditioned by a series of front-end electronics **82**. These conditioning circuits **82** may include various amplifiers, filters, and analog-to-digital converters. The digital signals output by front-end electronics **82** are then processed by a coincidence processor **84** to match gamma ray detections as potential coincidence events. When two gamma rays strike detectors approximately opposite one another, it is possible, absent the interactions of random noise and signal gamma ray detections, that a positron annihilation took place somewhere along the line between the detectors. Thus, the coincidences determined by coincidence processor **84** are sorted into true coincidence events and are ultimately integrated by data sorter **86**. The coincidence event data, or PET data, from data sorter **86** is received by the system control **50** at a PET data

receive port **88** and stored in memory **48** for subsequent processing **78**. PET images may then be reconstructed by image processor **44** and may be combined with MR images to produce hybrid structural and metabolic or functional images. Front-end electronics **82**, coincidence processor **84**, and data sorter **86** may each be external of scanner **58** or system control **50**, or may be integrated therein.

A blanking control or monitor **90** is also included in system control **50**. Blanking monitor **90** identifies and records times during which MR components **66-72** are active or transmitting. Blanking monitor **90** may use this timing data to gate PET data acquisition by detector **80** or signal conditioning by front-end electronics **82**, or may output a timing sequence to be applied during data processing by coincidence processor **84**, data sorter **86**, data processor **78**, or image reconstructor **44**.

Referring now to FIG. 3, a PET/CT imaging system **92**, useable with multi-modality imaging subsystem **20** of FIG. 1 is shown. It should be noted that in some embodiments, PET/CT imaging system **92** acquires CT data prior to obtaining the PET data. However, one skilled in the art will recognize that the PET and CT data may be acquired in different orders and combinations thereof (e.g., in an interleaved manner).

PET/CT imaging system **92** generally includes a gantry **94**, a patient table **96**, and a processing and control system **98** including a user input **100** with a display **102**. The gantry **94** provides mechanical support for imaging devices such as, for example, detectors, scanners, and transmitters that are used for scanning a patient **104**. The gantry **94** houses imaging devices such as, for example, PET detectors or x-ray detectors. It should be noted that the PET system may be a stationary annular detector or optionally may include a pin source.

In some embodiments, gantry **94** includes a plurality of PET detectors that are fixed and spaced on the gantry **94**, which are positioned radially outward from the axis. In accordance with other embodiments, the gantry **94** includes a plurality of detectors that are rotatable about the axis. For CT imaging, a rotating detectors and a source, for example, an x-ray tube **106** may be provided and optionally including a stationary detector ring for CT imaging may be provided. In other embodiments, a separate imaging gantry is provided for CT imaging.

The imaging devices on the gantry **94** acquire image data by scanning the patient **104** lying on the patient table **96**. The patient table **96** lies along the axis of the gantry **94**, and can be moved along this axis. Moving the patient table **96** enables the scanning of various portions of the patient **104**. The patient table **96** can be positioned at various axial positions along the axis.

The processing and control system **98** controls the positioning of the patient table **96**, and receives image data collected during scanning. In various embodiments, the processing and control system **98** controls the PET/CT imaging system **92** to acquire both emission and transmission image data of a volume of interest. For example, annihilation events may be detected as emission data, as well as transmission data from signal transmitted by a transmission source, such as the x-ray tube **106**, which pass through the volume of interest of the patient **104**. The transmission signals may get attenuated when the signals pass through the volume of interest and the detectors may collect data that is attenuated after the transmission signals pass through the patient **104**.

Various processors, sorters, and databases are used to acquire and manipulate emission and transmission data, which is used in accordance with various embodiments. The processors, sorters, and databases of FIG. 3 include acquisi-

tion circuitry **108**, an acquisition processor **110**, a transmission data database **112**, an emission data database **114**, and an image reconstruction processor **116**. The acquisition processor **110** is programmed to acquire emission data and generate an image based on the emission data acquired. Other computing components also may be included.

In some embodiments, an energy sorter **118** provides, for example, time, location, and energy data to a PET processor **120**. The PET processor **120** generally uses the received data to identify pairs of data, also known as coincidence pairs, coincident pair lines, and lines of response, corresponding to annihilation events that occurred inside the region of interest. After acquisition processor **110** identifies an annihilation event, the acquisition processor **110** updates data in the emission data database **114** to store information relating to the annihilation event. CT data is also stored in the transmission data database **112** based on transmission signals that pass through the patient **104** and are detected.

Thus, after an acquisition session has been completed and sets of transmission and emission data have been stored in databases **112** and **114**, respectively, image reconstruction processor **116** accesses the data in the databases **112** and **114** and uses the accessed data to generate images that may be requested by a system operator. Additionally, the sets of transmission and emission data are used by a scatter correction estimator module **122** to estimate scatter for the emission data based on both the transmission and emission data.

FIG. **4** sets forth a technique **200** for quantifying bone lesions in a patient using multi-modality three dimensional image data. According to various embodiments, technique **200** is performed by a computer processor such as, for example, CPU **12** of FIG. **1**. Technique **200** begins with step **202**, by selecting and loading one or more exams to be processed by the computer system corresponding to one or more imaging sessions during which multi-modality image data was acquired from a patient of interest. The multi-modality image data includes at least one set of anatomical image data acquired from the patient using an anatomical imaging modality, such as CT or MR for example, and at least one set of functional image data acquired from the patient using a functional imaging modality, such as PET or SPECT as examples. In the exemplary embodiment described below wherein the multi-modality image data includes three-dimensional CT and PET data, one CT series and one PET series from one of the loaded exams is selected in step **204**. In alternative embodiments, one MR series and one PET series, one CT series and one SPECT series, and/or one MR series and one SPECT series may be selected from the loaded exams in step **204**.

In optional step **206** (shown in phantom) technique **200** performs a registration of the image data of the CT and PET series in order to accurately overlay the anatomical and functional image data. Alternatively, in embodiments where the multi-modality image data is acquired from a combined PET/CT or a combined PET/MR imaging system, the acquired multi-modality image data may be automatically registered.

During step **208** an automatic bone segmentation is performed on the anatomical image data (e.g., data corresponding to the CT volume) using a bone segmentation subroutine **210** described in detail below with respect to FIG. **5**. In one embodiment, a patient skeletal image representing the skeleton of the patient is generated from a subset of the anatomical image data during step **208** based on the automatic bone segmentation.

Referring to FIG. **5**, the bone segmentation subroutine **210** of technique **200** is described in detail. As shown, bone segmentation subroutine **210** begins at step **212** by accessing the

anatomical image series, such as a CT series, selected at step **204** of technique **200**. In optional step **214** (shown in phantom) one or more views of the accessed image series is displayed on a user interface, such as on workstation **28** of FIG. **1**, to permit a user to review the image series and adjust the segmentation parameters. In step **216** bone segmentation subroutine **210** performs an automated segmentation to identify and define the skeletal structure within the anatomical image series using predefined segmentation parameters and/or the adjusted segmentation parameters defined in step **214**.

In embodiments where the anatomical image series includes CT data, bone segmentation subroutine **210** uses a thresholding technique to segment the image data. The thresholding technique may, for example, define image data having attenuation values within a predetermined range as corresponding to skeletal or bone material. In embodiments where the anatomical image series includes MR data, bone segmentation subroutine **210** segments the image data based on general CT anatomy atlas mapping or using ultrashort echo time sequences generated MR data, where with a thresholding algorithm the skeleton of the bone structure can be obtained. While various segmenting techniques are described herein, one having skill in the art will recognize that embodiments of the invention may segment different anatomical region bone structures using alternative techniques, such as, for example, clustering, level set, and region growing.

After performing the automated segmentation, bone segmentation subroutine **210** fills the cavities on the defined bone structure to add the bone marrow, the spinal-cord, and the brain in step **218**. In step **220** the jaw-bone is detected based on the bone/soft tissue ratio in order to eventually remove the brain. Bone segmentation subroutine **210** uses the detected jaw-bone to separate the whole head and segment the brain in step **222**. The segmented brain is removed from the anatomical image data in step **224**. Bone segmentation subroutine **210** then generates a patient skeleton image from a subset of the anatomical image data representing the bone of the patient and displays the image in one or more views at step **226** for user review.

If the segmentation is deemed acceptable **228** at decision block **230**, the bone segmentation subroutine **210** outputs the segmented skeletal structure of the patient and ends at block **232**. If the segmentation is not acceptable **234**, bone segmentation subroutine **210** proceeds to decision block **236** to determine whether the segmentation can be corrected by manual editing. If the segmentation cannot be corrected by manual editing **238**, bone segmentation subroutine **210** returns to step **214** to adjust the segmentation parameters and restart the segmentation process. If the segmentation can be corrected by manual editing **240**, bone segmentation subroutine **210** proceeds to step **242**, wherein a user manually edits the current results of the segmentation, such as by modifying the contours of the resulting skeletal structure. After the manual editing procedure is complete, bone segmentation subroutine **210** returns to decision block **230** to determine whether the result of the edited segmentation is acceptable.

Referring again to FIG. **4**, after the bone segmentation subroutine **210** of step **208** is complete, technique **200** continues to step **244** wherein technique **200** uses the segmented anatomical image data output from bone segmentation subroutine **210** and the functional image data selected at step **204** to detect bone lesions at using a lesion detection subroutine **246**, as described in detail in FIG. **6**.

Referring now to FIG. **6**, the lesion detection subroutine **246** begins at step **248** by accessing the functional image data set corresponding to the image series selected at step **204** of technique **200**. In step **250** lesion detection subroutine **246**

automatically identifies possible lesion candidates in the functional image data set using a thresholding algorithm. In one embodiment, the possible lesion candidates are identified based on the standardized uptake values (SUV) of image voxels within the PET data series. For example, lesion candidates may be defined as regions of voxels having a mean SUV within a predetermined range of values. During step 250, lesion detection subroutine 246 also assigns bookmarks to the lesion candidates. According to various embodiments, the assigned bookmarks may be automatically assigned to lesion candidates by a computer processor, such as CPU 12 of FIG. 1, and/or manually assigned by a user. The bookmarks may include identifying information for the lesion candidates including for example, a unique lesion identifier and/or the positional coordinates for the respective lesion candidate.

Images of the lesion candidates are generated in step 252 and displayed as an overlay on the patient skeletal image generated from the segmented skeletal structure output from bone segmentation subroutine 210 of FIG. 5. In one embodiment, the segmented skeletal structure from a CT image series is projected into a PET image that includes the lesion candidates. In decision block 254, the results of the automated lesion candidate detection step are reviewed to determine if the result of the lesion candidate detection is acceptable. If the lesion detection is not acceptable 256, lesion detection subroutine 246 proceeds to step 258 in which the lesion detection parameters are adjusted. However, if the lesion detection is acceptable 260, lesion detection subroutine 246 is prompted to move on to step 262 and classify the identified lesion candidates as being either bone lesions (i.e., lesions located within the skeletal structure) or non-bone lesions (i.e., lesions not located in the skeletal structure).

While step 252 is described above as occurring prior to decision block 254, lesion detection subroutine 246 may perform step 252 after decision block 254 determines that the results of the lesion candidate identification is acceptable in alternative embodiments.

To assist with classifying lesion candidates as being either bone lesions or non-bone lesions at step 262, lesion detection subroutine 246 uses the segmented CT volume of step 208. In particular, lesion detection subroutine 246 compares the location of the lesion candidates to the location of the identified skeletal structure. Any lesion candidates that fall within the location of the skeletal structure are automatically identified as bone lesions and any lesion candidates that are outside location of the skeletal structure are automatically identified as non-bone lesions. Because lesion detection subroutine 246 automatically classifies lesions as being bone or non-bone based on the identified location of the lesion candidate with respect to the identified location of the patient's skeletal structure, lesion candidates may be correctly classified as being bone lesions with high accuracy. The bookmark information for the lesion candidates may be updated following lesion detection subroutine 246 to include information to indicate whether the lesion candidate is a bone lesion or a non-bone lesion. Further, data corresponding to the classified bone lesions may be combined and interpreted as a common object for quantitative analysis, rather than reviewing and determining quantitative data for each lesion separately. Lesion detection subroutine 246 ends at block 264.

Referring again to FIG. 4, the results of the lesion classification determined during lesion detection subroutine 246, as well as the bookmark information for the lesion candidates, are optionally output to a database for storage at step 266 (shown in phantom). Technique 200 also uses the results of the lesion classification in step 268 to calculate quantitative information regarding the skeletal structure and detected

bone lesions of the patient. It is contemplated that quantitative information regarding the skeletal structure and detected bone lesions may be in regard to individual bone lesions or the total of all bone lesions, according to various embodiments.

The quantitative information is displayed at step 270.

According to one embodiment, technique 200 calculates a patient skeletal metric that represents a total skeletal volume of the patient, a bone lesion metric that represents a total bone lesion volume of the patient, and a bone lesion index or lesion burden at step 268. The total skeletal volume is a quantitative value calculated based on the final bone segmentation output from bone segmentation subroutine 210 and represents the total volume of bone within the patient. The total bone lesion volume is a quantitative value calculated based on a total bone volume of the lesion candidates classified as bone lesions by lesion detection subroutine 246. According to one embodiment, the bone lesion index is calculated as a ratio of the total bone lesion volume to the total skeletal volume and is represented as a percentage. The bone lesion index quantifies the amount of disease in the patient's skeletal structure and represents a total tumor burden of the skeletal structure.

In some embodiments, portions of the selected anatomical and functional image series, visual representations of the lesion candidates, and the quantitative information calculated by technique 200 may be displayed on a graphical user interface (GUI) 272 as illustrated in FIG. 7. As shown, GUI 272 includes a menu display region 274, which contains an anatomical image selection menu 276, a functional image selection menu 278, and various command buttons 280. Anatomical image selection menu 276 and functional image selection menu 278 allow the user to select which multi-modality image series to load and analyze. According to various embodiments, command buttons 280 allow the user to interact with select steps of technique 200 and subroutines 210, 246. For example, one command button 280 may permit a user to adjust a segmentation parameter of segmentation subroutine 210 at step 214 or adjust lesion detection parameters of lesion detection subroutine 246 at step 258.

According to an exemplary embodiment, GUI 272 includes an image display region 282 that is separated into multiple windows in order to display multiple images. For example, image display 282 may include a first window 284 that displays a CT image, a second window 286 that displays a corresponding PET image, and a third window 288 that displays the CT image overlaid on the PET image. Fourth, fifth, and sixth windows 290, 292, 294 include various views of the segmented skeletal volume obtained from the CT image using bone segmentation subroutine 210 overlaid with contours of lesion candidates 296, 298 detected using lesion detection subroutine 246. In alternative embodiments, image display region 282 includes one or more selected images corresponding to the multi-modality image series and the detected lesions. Image display region 282 may further be configured to display bone lesions and non-bone lesions in different colors or with different shading to visually distinguish detected bone lesions from non-bone lesions. As one example, a lesion candidate classified as a bone lesion, such as lesion candidate 296, may be displayed with a red contour, and a lesion candidate classified as a non-bone lesion, such as lesion candidate 298, may be displayed with a blue contour. According to various embodiments, image display region 282 may be configured as an interactive workspace to permit a user to interact with the images to perform various tasks including, for example, manually identify lesion candidates, delete an automatically identified lesion, and/or change the classification of an automatically detected lesion, as non-limiting examples.

Analysis display region **300** includes a visual display of the quantitative or statistical data computed in step **268** of technique **200**. This data can include the total skeleton volume of the patient, the total volume of bone lesions and non-bone lesions, the total bone lesion volume (i.e., total tumor burden), the calculated bone lesion index, and other statistical information calculated during step **268**. An information display region **302** is also located within in GUI **272** to provide the user with patient and scan information including, for example, name, age, the date of scans, and/or various parameters associated with the scans.

GUI **272** also includes a bookmark display region **304** that lists identifying information for the lesion candidates flagged by either the computer system or the user. According to various embodiments, bookmark display region **304** may include a unique identifier for a particular lesion candidate, the coordinates for that particular lesion candidate, and whether the particular lesion candidate is a bone-lesion or a non-bone lesion.

Optionally, one or more of regions **274**, **282**, **300**, **302**, and **304** may be configured as a control panel to permit a user to input and/or select data through input fields, dropdown menus, etc. It is noted that the arrangement of GUI **272** is provided merely for explanatory purposes, and that other GUI arrangements, field names, and visual outputs may take different forms within the scope of various embodiments of the invention. Additional display techniques may also include temperature gauges, graphs, dials, font variations, annotations, and the like.

A technical contribution for the disclosed method and apparatus is that it provides for a computer implemented system and method of detecting bone lesions using multi-modality image data.

One skilled in the art will appreciate that embodiments of the invention may be interfaced to and controlled by a computer readable storage medium having stored thereon a computer program. The computer readable storage medium includes a plurality of components such as one or more of electronic components, hardware components, and/or computer software components. These components may include one or more computer readable storage media that generally stores instructions such as software, firmware and/or assembly language for performing one or more portions of one or more implementations or embodiments of a sequence. These computer readable storage media are generally non-transitory and/or tangible. Examples of such a computer readable storage medium include a recordable data storage medium of a computer and/or storage device. The computer readable storage media may employ, for example, one or more of a magnetic, electrical, optical, biological, and/or atomic data storage medium. Further, such media may take the form of, for example, floppy disks, magnetic tapes, CD-ROMs, DVD-ROMs, hard disk drives, and/or electronic memory. Other forms of non-transitory and/or tangible computer readable storage media not list may be employed with embodiments of the invention.

A number of such components can be combined or divided in an implementation of a system. Further, such components may include a set and/or series of computer instructions written in or implemented with any of a number of programming languages, as will be appreciated by those skilled in the art. In addition, other forms of computer readable media such as a carrier wave may be employed to embody a computer data signal representing a sequence of instructions that when executed by one or more computers causes the one or more computers to perform one or more portions of one or more implementations or embodiments of a sequence.

In accordance with one embodiment of the present invention, a non-transitory computer readable storage medium has stored thereon a computer program and represents a set of instructions that when executed by the computer causes the computer to access a first set of data of a first data type and a second set of data of a second data type. The first set of data is acquired from a patient with a first imaging modality, and the second set of data is acquired from the patient with a second imaging modality. The set of instructions also causes the computer to perform a segmentation on the first set of data to identify a subset of the first set of data corresponding to a skeletal structure of the patient. Further, the set of instructions causes the computer to automatically calculate a patient skeletal metric from the subset of the first set of data, the patient skeletal metric representing a total bone volume of the patient. In addition, the set of instructions causes the computer to detect at least one lesion in the second set of data, classify the at least one lesion as a one of a bone lesion and a non-bone lesion, and automatically calculate a bone lesion metric based on the classification. The set of instructions further causes the computer to calculate a lesion burden as a ratio of the bone lesion metric and the patient skeletal metric.

According to another embodiment of the invention, a method includes accessing an anatomical image data set acquired from a patient and accessing a function image data set acquired from the patient. The method also includes identifying a subset of the anatomical image dataset corresponding to bone and calculating a skeletal volume from the subset of the anatomical image dataset. In addition, the method includes identifying a set of lesions from the function image dataset, comparing the set of lesions to the subset of the anatomical image dataset to identify at least one bone lesion within the set of lesions, and automatically calculating a bone lesion burden based on the at least one bone lesion. Further, the method includes calculating a bone lesion index from the bone lesion burden and the skeletal volume.

According to yet another embodiment of the invention, a medical diagnostic tool includes an image acquisition system configured to acquire multi-modality image data, a first database having stored thereon a first image dataset acquired from a patient using a first imaging modality, a second database having stored thereon a second image dataset acquired from the patient using a second imaging modality, and a computer processor. The computer processor is programmed to access the first and second image datasets. The computer processor is further programmed to segment the first image dataset to define a set of skeletal data, identify a location of a lesion candidate in the second image set, compare the location of the lesion candidate to the set of skeletal data, and classify the lesion candidate as one of a bone lesion and anon-bone lesion based on the comparison to define a set of bone lesion data. In addition, the computer processor is programmed to calculate a lesion burden based on a ratio of the set of bone lesion data and the set of skeletal data. The medical diagnostic tool also includes a graphical user interface (GUI) constructed to display an image generated from overlaying a first image corresponding to the set of skeletal data on a second image corresponding to the set of bone lesion data.

This written description uses examples to disclose the invention, including the best mode, and also to enable any person skilled in the art to practice the invention, including making and using any devices or systems and performing any incorporated methods. The patentable scope of the invention is defined by the claims, and may include other examples that occur to those skilled in the art. Such other examples are intended to be within the scope of the claims if they have structural elements that do not differ from the literal language

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of the claims, or if they include equivalent structural elements with insubstantial differences from the literal languages of the claims.

What is claimed is:

1. A non-transitory computer readable storage medium having a computer program stored thereon and representing a set of instructions that, when executed by a computer, causes the computer to:

access a first set of data of a first data type, the first set of data acquired from a patient with a first imaging modality;

access a second set of data of a second data type, the second set of data acquired from the patient with a second imaging modality;

perform a segmentation on the first set of data to identify a subset of the first set of data corresponding to a skeletal structure of the patient, the segmentation including:

detect a jaw-bone of the patient in the subset of the first set of data; and

remove data corresponding to a patient brain from the subset of the first set of data based on the detected jaw-bone;

generate a segmented skeletal structure of the patient by filling a plurality of cavities in the subset of the first set of data, the plurality of cavities corresponding to bone marrow locations within the skeletal structure of the patient;

automatically calculate a patient skeletal metric from the segmented skeletal structure, the patient skeletal metric representing a total bone volume of the patient;

detect at least one lesion in the second set of data;

classify the at least one lesion as one of a bone lesion and a non-bone lesion;

automatically calculate a bone lesion metric based on the classification; and

calculate a lesion burden as a ratio of the bone lesion metric and the patient skeletal metric.

2. The computer readable storage medium of claim 1 wherein the set of instructions further causes the computer to: automatically calculate a total volume of lesions classified as bone lesions; and use the calculated total volume to calculate the bone lesion metric.

3. The computer readable storage medium of claim 1 wherein the set of instructions further causes the computer to: generate an image of the skeletal structure of the patient from the skeletal structure; generate an image of the at least one lesion from the second set of data; and overlay the image of the at least one lesion on the image of the skeletal structure of the patient.

4. The computer readable storage medium of claim 1 wherein the set of instructions further causes the computer to register the first set of data with the second set of data.

5. The computer readable storage medium of claim 1 wherein the first imaging modality comprises a computed tomography (CT) imaging system; and

wherein the second imaging modality comprises a positron emission tomography (PET) imaging system.

6. The computer readable storage medium of claim 1 wherein the first imaging modality comprises an magnetic resonance (MR) imaging system; and

wherein the second imaging modality comprises a positron emission tomography (PET) imaging system.

7. The computer readable storage medium of claim 1 wherein the set of instructions further causes the computer to

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automatically assign a bookmark to the at least one lesion, the bookmark comprising identifying information for the at least one lesion.

8. A method comprising:

accessing an anatomical image data set acquired from a patient;

accessing a functional image data set acquired from the patient;

identifying a subset of the anatomical image dataset corresponding to bone using a segmentation subroutine, the segmentation subroutine having bone segmentation and cavity-filling wherein cavities within bone structure are filled to add image data corresponding to at least one of bone marrow, spinal cord, and brain, wherein the bone segmentation includes:

detecting a jaw-bone of the patient in the subset of the anatomical image dataset; and

removing data corresponding to the patient's brain based on the detected jaw-bone;

calculating a skeletal volume from the subset of the anatomical image dataset;

identifying a set of lesions from the functional image dataset;

comparing the set of lesions to the subset of the anatomical image dataset to identify at least one bone lesion within the set of lesions;

automatically calculating a bone lesion burden based on the at least one bone lesion; and

calculating a bone lesion index from the bone lesion burden and the skeletal volume.

9. The method of claim 8 further comprising:

acquiring the anatomical image dataset from one of a computed tomography (CT) imaging system and a magnetic resonance (MR) imaging system; and

acquiring the functional image dataset from one of a photon emission tomography (PET) imaging system and a single-photon emission computed tomography (SPECT) imaging system.

10. The method of claim 8 further comprising:

generating a patient skeleton image from the subset of the anatomical image dataset, wherein the patient skeleton image represents a patient skeleton;

generating at least one lesion image from the identification of the at least one bone lesion;

displaying the at least one lesion image as an overlay on the patient skeleton image on a graphical user interface (GUI); and

displaying at least one of the bone lesion burden and the bone lesion index on the GUI.

11. The method of claim 8 further comprising:

defining a first subset of the set of lesions as bone lesions; and

defining a second subset of the set of lesions as non-bone lesions.

12. The method of claim 8 further comprising manually modifying at least one of a parameter of the segmentation subroutine and a parameter of a lesion detection subroutine used to identify the set of lesions from the functional image dataset.

13. The method of claim 8 further comprising assigning bookmarks to the set of lesions.

14. A medical diagnostic tool comprising:

a computer processor programmed to:

access first and second image datasets;

segment the first image dataset to define a set of skeletal data;

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perform a bone segmentation subroutine on the set of skeletal data to define a modified set of skeletal data, the bone segmentation subroutine comprising:
 detecting a jaw-bone of the patient in the set of skeletal data; and
 removing data corresponding to a brain from the set of skeletal data based on the detected jaw-bone;
 identify a location of a lesion candidate in the second image dataset;
 compare the location of the lesion candidate to the modified set of skeletal data;
 classify the lesion candidate as one of a bone lesion and a non-bone lesion based on the comparison to define a set of bone lesion data;
 calculate a lesion burden based on a ratio of the set of bone lesion data and the modified set of skeletal data;
 and
 a graphical user interface (GUI) constructed to display an image generated from overlaying a first image corresponding to the modified set of skeletal data on a second image corresponding to the set of bone lesion data.

15. The medical diagnostic tool of claim **14** wherein the first image dataset is acquired from a first imaging modality, wherein the first imaging modality is one of computed tomography (CT) imaging system and magnetic resonance (MR) imaging system; and

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wherein the second image dataset is acquired from a second imaging modality, wherein the second imaging modality is one of positron emission tomography (PET) imaging system and single-photon emission computed tomography (SPECT) imaging system.

16. The medical diagnostic tool of claim **14** wherein the GUI comprises:
 an analysis display window constructed to display the lesion burden as a quantitative value; and
 a user input region constructed to allow a user to modify at least one parameter of the segmentation.

17. The medical diagnostic tool of claim **14** wherein the computer processor is further programmed to:
 identify a plurality of lesion candidates;
 classify a subset of the plurality of lesion candidates as bone lesions;
 calculate a volume of the bone lesions; and
 define the lesion burden from the volume of bone lesions.

18. The medical diagnostic tool of claim **14** wherein in performing the bone segmentation subroutine the computer processor is further programmed to fill cavities within the set of skeletal data corresponding to bone marrow.

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